II. CLAIMS

- 1. (Currently amended) A $G_{\text{oq-Costducin}}$ chimeric G-protein wherein the last 44 amino acids of the G_{oq} protein sequence are replaced with a 44 amino acid unit of Gustducin, where such 44 amino acid unit of Gustducin is the last 44 amino acids of SEQ ID NO:2.
- 2. (Previously Presented) The chimeric $G_{\alpha q}$ -Gustducin according to elaim 1 claim 18 characterised in that it is a $G_{\alpha 15~{\rm or}~16\text{-}Qustducin}$ protein.

3-5. Cancelled

- 6. (Currently Amended) A G-protein according to claim 1 encoded for by the nucleic acid set forth in SEQ ID NO:1.
- 7. (Currently Amended) A nucleic acid comprising the nucleotide sequence set forth in SEQ ID $\underline{\text{NO:}}1$ encoding for a G-protein according to claim 1.
- 8. (Currently Amended) An expression vector comprising nucleic acid comprising the nucleotide sequence set forth in SEQ ID $\underline{\text{NO:}}1$ encoding for a G-protein according to claim 1.
- 9. (Previously Presented) A host cell transformed with an expression vector according to claim 8.
- 10. (Previously Presented) A method of producing a chimeric Gprotein according to claim 1 comprising the step of culturing
 host cells having contained therein an expression vector
 encoding for the chimeric G-protein, under conditions sufficient

for expression of said G-protein, thereby causing production of the protein, and recovering the protein produced by the cell.

- 11. (Previously Presented) A method of analysis and discovery of modulators of bitter taste receptors using the chimeric proteins according to defined in claim 1.
- 12. (Previously Presented) A method according to claim 11 employing a mammalian cell-based assay employing a transfected gene or cDNA encoding a chimeric protein of the invention and a taste receptor, the method comprising the steps of contacting a compound with cells, and determining the functional effect of the compound on chimeric G-protein.
- 13. Previously Presented) A method according to claim 10 wherein the functional effect is determined by measuring the changes in intracellular messengers IP3 or calcium²⁺.

14-17. Cancelled

- 18. (New) A $G_{\text{aq-Gustducin}}$ chimeric G-protein wherein the last 44 amino acids of the G_{aq} protein sequence are replaced with a 44 amino acid unit of Gustducin, where such 44 amino acid unit of Gustducin is the last 44 amino acids of SEQ ID NO:2, and wherein the resulting $G_{\text{aq-gust44}}$ chimeric G-protein has a sequence homology of at least 80% in the last 44 amino acids of SEQ ID NO:2.
- 19. (New) The chimeric G-protein of claim 18 having a sequence homology of at least 90% in the last 44 amino acids of SEQ ID NO:2.

- 20. (New) The chimeric G-protein of claim 18 having a sequence homology of at least 95% in the last 44 amino acids of SEQ ID No:2.
- 21. (New) A $G_{\text{aq-Guntducin}}$ chimeric G-protein wherein the last 44 amino acids of the G_{aq} protein sequence are replaced with a 44 amino acid unit of Gustducin, where such 44 amino acid unit of Gustducin is the last 44 amino acids of SEQ ID NO:2, and wherein the resulting $G_{\text{aq-gunt44}}$ chimeric G-protein has a sequence homology of at least 80% to SEO ID NO:2.
- 22. (New) The chimeric G-protein of claim 21 having a sequence homology of at least 90% to SEO ID NO:2.
- 23. (New) The chimeric G-protein of claim 21 having a sequence homology of at least 95% to SEQ ID NO:2.
- 24. (New) A $G_{\text{eq-Gustducin}}$ chimeric G-protein wherein the last 44 amino acids of the G_{eq} protein sequence are replaced with a 44 amino acid unit of Gustducin, where such 44 amino acid unit of Gustducin is the last 44 amino acids of SEQ ID NO:2, and wherein the resulting $G_{\text{eq-gust44}}$ chimeric G-protein has a sequence homology of at least 80% to SEQ ID NO:2 and the chimeric protein binds to one or more of the human bitter, sweet and umami taste receptors.
- 25. (New) The chimeric G-protein of claim 24 having a sequence homology of at least 90% to SEQ ID NO:2.
- 26. (New) The chimeric G-protein of claim 24 having a sequence

homology of at least 95% to SEQ ID NO:2.

- 27. (New) The chimeric $G_{\rm eq}\text{-}Gustducin$ according to claim 18 characterised in that it is a $G_{\rm cd5~or~16-Gustducin}$ protein.
- 28. (New) A nucleic acid encoding for a G-protein according to claim 18.
- 29. (New) An expression vector comprising nucleic acid comprising the nucleotide sequence encoding for a G-protein according to claim 18.
- 30. (New) A host cell transformed with an expression vector according to claim 29.
- 31. (New) A method of producing a chimeric G-protein according to claim 18 comprising the step of culturing host cells having contained therein an expression vector encoding for the chimeric G-protein, under conditions sufficient for expression of said G-protein, thereby causing production of the protein, and recovering the protein produced by the cell.
- 32. (New) A method of analysis and discovery of modulators of bitter taste receptors using the chimeric proteins according to defined in claim 18.
- 33. (New) A method according to claim 32 employing a mammalian cell-based assay employing a transfected gene or cDNA encoding a chimeric protein of the invention and a taste receptor, the method comprising the steps of contacting a compound with cells, and determining the functional effect of the compound on

chimeric G-protein.

34. (New)) A method according to claim 31 wherein the functional effect is determined by measuring the changes in intracellular messengers IP3 or calcium²⁺.